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The correction of reflection intensities for incomplete absorption of high-energy X-rays in the CCD phosphor

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It is shown that incomplete absorption of the X-ray beam in the phosphor of an area detector causes an incident-angle dependence of the recorded X-ray intensities. An energy scan of a SMART-6000 CCD (charge-coupled device) phosphor using synchrotron radiation shows the correction to be of importance above about 17 keV. Intensities of single reflections, each collected several times at different angles of incidence on the phosphor surface, show a pronounced angle-dependence at shorter wavelengths. Both conventional structural refinement and multipole charge density studies confirm that an oblique-incidence correction leads to improved quality of the results. Atomic displacement parameters will be systematically biased when the correction is not applied. For a $\lambda = 0.394 \text{ \AA}$ data set, neglecting the correction gives rise to artifacts in the deformation density maps that are likely to lead to misinterpretation of the experimental results.

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1. Introduction

CCD detectors are now widely used in both macromolecular and small-molecule crystallographic studies. While in macromolecular crystallography radiation with wavelengths above 1 \AA is commonly used, in small-molecule structure analysis, and particularly in charge density studies, Mo $K\alpha$ or shorter wavelength X-rays have distinct advantages and are widely employed. For such short wavelengths, the phosphor layer of the detector does not completely absorb the X-ray beam.

As was pointed out by Gruner (1993), but not always taken into account, when absorption is incomplete the phosphorescence will depend on the path length of the beam in the phosphor, and therefore on the angle of incidence α , defined as the angle between the diffracted beam and the normal to the CCD surface. This is the 'thin phosphor regime'. On the other hand, in the 'thick phosphor regime', where absorption of the X-ray beam in the phosphor is very close to 100%, as typically occurs for softer radiation, the efficiency of light transmission through the phosphor layer becomes the dominant effect. In the 'thick phosphor regime', less intensity may actually be recorded at larger angles of incidence, for which the absorption occurs closer to the entry surface of the phosphor (Gruner, 1993).

For incomplete absorption, the path length of the X-ray beam in the phosphor equals $t/\cos(\alpha)$, where t is the thickness of the phosphor. Thus, while for perpendicular incidence the absorption is $(1 - T_{\perp})$, where T_{\perp} is the transmission at perpendicular incidence, it is $\{1 - \exp[\ln(T_{\perp})/\cos(\alpha)]\}$ for a beam with incident angle α . In earlier work with imaging plates (Zaleski *et al.*, 1998) and short-wavelength radiation,

the observed intensity I_{obs} was reduced to an equivalent perpendicular-incidence intensity I_{\perp} with the expression

$$I_{\perp} = I_{\text{obs}}(1 - T_{\perp})/\{1 - \exp[\ln(T_{\perp})/\cos(\alpha)]\}. \quad (1)$$

This oblique-incidence correction proved to correct satisfactorily quite pronounced differences between the intensities of the same reflection recorded at different angles of incidence on BASF III imaging plates.

We show here that a similar correction is required for short-wavelength data collected with a SMART-6000 CCD detector using a $\text{Gd}_2\text{O}_2\text{S}$ phosphor with a surface density of 50 mg cm^{-2} , and that expression (1) leads to significant improvements in the experimental data sets. No other corrections were tested in this study.

2. Experimental

A $25 \times 25 \text{ cm}$ piece of a $\text{Gd}_2\text{O}_2\text{S}$ phosphor with a surface density of 50 mg cm^{-2} and thickness of 0.2 mm , designed for the SMART-6000 CCD, was provided by Bruker AXS Inc. Its perpendicular transmission was measured at the SUNY X3 beamline at the National Synchrotron Light Source at Brookhaven National Laboratory, using both 19.3 keV (0.643 \AA) and 31.45 keV (0.394 \AA) radiation from sideways-mounted Si(111) and Si(220) crystals, respectively. To eliminate high-order contamination, the $\bar{3}\bar{1}\bar{2}$ reflection from a crystal of diaquabis(hydrogen phthalato)copper(II) (Bartl & Kuppers, 1980; Rodrigues *et al.*, 2002), rather than the primary beam, was used. The perpendicular transmission was measured as 0.1237 and 0.5610 at the two wavelengths,

Table 1

Comparison of R_{merge} and final agreement factors with and without correction of the data for oblique incidence.

	2θ CCD ($^{\circ}$)	λ (\AA)	$(\sin \theta/\lambda)_{\text{max}}$ (\AA^{-1})	T_{\perp}	α ($^{\circ}$) \dagger	R_{merge} (no correction)	R_{merge} (with correction)	$R_{\text{refinement}}$ (no correction)	$R_{\text{refinement}}$ (with correction)
$\text{CuC}_{28}\text{H}_{24}\text{BN}_2\text{F}_4\text{S}_2\ddagger$	20	0.643	0.89	0.1237	0.33–62.18	6.01%	5.48%	3.59%	3.37%
$\text{ZrC}_{17}\text{H}_{31}\text{P}\ddagger$	25	0.643	0.97	0.1237	0.45–68.16	7.05%	6.70%	4.35%	4.24%
$\text{Cu}(\text{C}_8\text{H}_5\text{O}_4)_2\cdot 2\text{H}_2\text{O}\ddagger$	30	0.394	1.55	0.561	0.13–61.02	5.85%	4.30%	5.03%	4.30%
$\text{Cu}(\text{C}_8\text{H}_5\text{O}_4)_2\cdot 2\text{H}_2\text{O}\S$	0	0.394	1.02	0.561	1.49–51.16	4.17%	4.06%	3.98%	3.90%
$\text{CuC}_8\text{H}_{14}\text{N}_2\text{O}_3\ddagger$	30	0.394	1.51	0.561	0.06–58.24	7.43%	5.27%	5.45%	4.22%
$\text{SrTiO}_3\S$	35	0.394	1.62	0.561	1.13–60.70	6.80%	6.06%	2.92%	2.05%
$\text{SrHfO}_3\S$	32	0.394	1.58	0.561	0.15–61.57	9.67%	8.51%	5.14%	4.27%

\dagger The angle between the diffracted beam and the vector normal to the CCD phosphor surface. \ddagger Measured at $T = 17$ K. \S Measured at $T = 293$ K.

respectively. The perpendicular transmission of the phosphor in the full 17–32 keV energy range was recorded at the ChemMat CARS beamline at the Advanced Photon Source at Argonne National Laboratory. The results are shown in Fig. 1.

To examine the incidence dependence of the intensities, two sets of measurements were made at the X3 beamline on crystals of 2-dimethylsulfuranylidene-1,3-indanedione (0.643 \AA) and diaquabis(hydrogen phthalato)copper(II) (0.394 \AA), with the detector positioned at $2\theta = 40, 30, 20, 10, 0$ and -10° . At each position, 110 frames with an oscillation range of 0.3° were collected. All intensities were integrated with the *SAINTE* (1999) program; no reflections were rejected. A vendor-supplied file for the floodfield correction, measured with Mo $K\alpha$ radiation and a 17.5 cm source-to-detector distance, was used. The intensities were corrected for the decay of the synchrotron beam using as reference the counts in an ionization chamber located between the beam defining slits and the SMART shutter.

Finally, as summarized in Table 1, the oblique-incidence correction was tested on several complete data sets collected at the X3A1 beamline. Equivalent and multiple measured reflections were averaged using the program *SORTAV* (Blessing, 1997). The program *SHELX* (Sheldrick, 1990) was used for structure refinement.

3. Analysis of the angle dependence

As shown in Fig. 1, for the phosphor tested the absorption at perpendicular incidence is not complete even at 17 keV, at

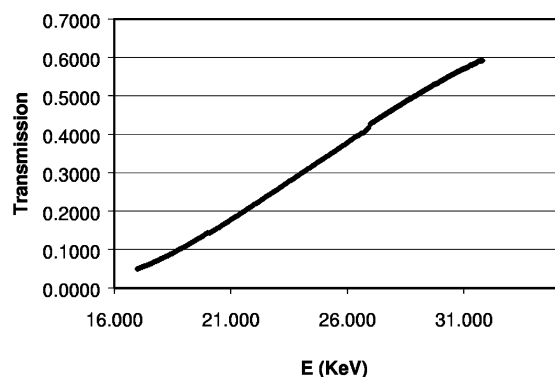


Figure 1
The transmission versus energy of the phosphor of the SMART 6000 CCD.

which the transmission is about 5%. This means that even with Mo $K\alpha$ radiation ($E = 17.45$ keV) the correction is desirable if an accuracy of better than 5% in the intensities is required, and a large detector positioned close to the crystal is employed. Temperature factors will be biased if the correction is ignored under such conditions. In the data collection procedure used with SMART detectors, a floodfield correction is applied to account for variation of the sensitivity across the

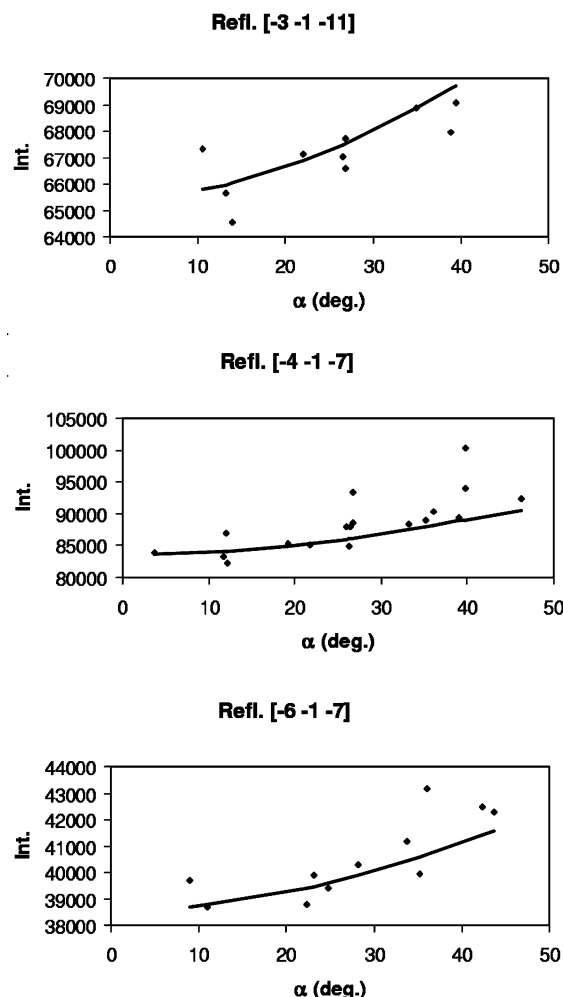


Figure 2
Observed (dot) and calculated (solid line) intensity versus angle of incidence for selected reflections of a $\text{C}_{11}\text{H}_{10}\text{O}_2\text{S}$ crystal collected at 0.643 \AA . The calculated values are from expression (1) with $T_{\perp} = 0.1237$.

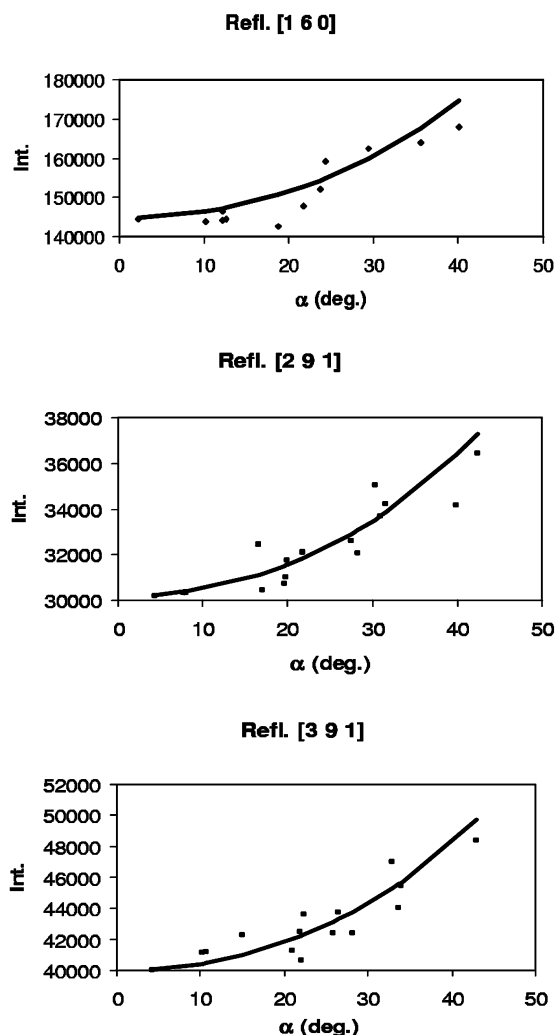


Figure 3 Observed (dot) and calculated (solid line) intensity *versus* angle of incidence for selected reflections of a $[\text{Cu}(\text{C}_8\text{H}_5\text{O}_4)_2(\text{H}_2\text{O})_2]$ crystal collected at 0.394 \AA . The calculated values are from expression (1) with $T_{\perp} = 0.561$.

detector surface using a pixel by pixel correction file, supplied with the detector or determined in a separate experiment. Such a correction would account for the differential absorption in the phosphor only if the same wavelength is employed and the source-to-detector distance is identical to the crystal-to-detector distance used in the diffraction experiment. These conditions are generally not fulfilled. At synchrotron sources the wavelength can be matched approximately by using an appropriate fluorescent foil as source, but this requires a much larger source-to-detector distance to avoid diffuse powder lines from the foil affecting the correction.

At the higher energies often used at synchrotron beamlines, the oblique-incidence correction becomes imperative. In accordance with the absorption curve of Fig. 1, the measurements at both 0.643 \AA and 0.394 \AA show a clear incident-angle dependence of the recorded intensities (Figs. 2 and 3). The full line represents the correction according to expression (1). As expected, the correction is most pronounced for the shorter wavelength.

Table 2

Comparisons of some equivalent isotropic temperature factors obtained without and with correction of the data for oblique incidence, $\lambda = 0394 \text{ \AA}$ and $T_{\perp} = 0561$.

	Atom	U_{eq} ($\times 10^3 \text{ \AA}^2$) (no correction)	U_{eq} ($\times 10^3 \text{ \AA}^2$) (with correction)
$[\text{Cu}(\text{C}_8\text{H}_5\text{O}_4)_2(\text{H}_2\text{O})_2]$ ($2\theta = 30^\circ$) [†]	Cu	3.94 (1)	4.85 (1)
$[\text{Cu}(\text{C}_8\text{H}_5\text{O}_4)_2(\text{H}_2\text{O})_2]$ ($2\theta = 0^\circ$) [‡]	Cu	23.87 (4)	26.24 (4)
$\text{CuC}_8\text{H}_{14}\text{N}_2\text{O}_3$ [†]	Cu	3.41 (1)	4.09 (1)
SrTiO_3 ($2\theta = 35^\circ$) [‡]	Sr	6.53 (7)	7.25 (5)
	Ti	4.40 (6)	5.07 (4)
	O	7.9 (1)	8.5 (1)
SrHfO_3 ($2\theta = 32^\circ$) [‡]	Sr	2.43 (2)	3.20 (2)
	Hf	11.88 (9)	12.23 (7)
	O(1)	61 (6)	83 (8)
	O(2)	13.0 (7)	13.4 (6)

[†] Measured at 17 K. [‡] Measured at 293 K.

The SMART-6000 CCD has a detector area of $92 \times 92 \text{ mm}$. At a crystal-to-detector distance of 43 mm , commonly used at the X3 beamline, the incidence angle α can in principle vary from 0 to 71.7° . With T_{\perp} equal to 0.561 (0.394 \AA), expression (1) predicts that the observed intensity at the maximal deviation from perpendicular incidence will be 1.92 times that of the same reflection recorded at normal incidence. Though the maximal angle is seldom reached, the correction can be very significant, as shown in Figs. 2 and 3. The correction is supported by the results shown, which indicate that expression (1) accounts reasonably well for the incident-angle dependence and allows reduction of all intensity measurements to the appropriate I_{\perp} value.

4. Effect of the correction

In Table 1, R_{merge} values and R factors from conventional structure refinements of a number of data sets collected at different wavelengths and different resolutions are tabulated before and after the oblique-incidence correction. In all cases both R_{merge} and the R factor from the least-squares refinement improve on application of the correction. The effect of the correction on R_{merge} may be especially large for data collected for charge density studies, in which several detector 2θ angles are often used. Thus, identical or equivalent reflections are very likely collected at different angles of incidence, leading to a significant deterioration in R_{merge} prior to correction.

The correction tends to decrease the intensity of the high-order reflections, which, given that the selection of the detector 2θ angle is optimized for efficient data collection, tend to occur at larger incident angles. Thus, the temperature factors are artificially lowered before the correction is applied. This is confirmed by the results shown in Table 2.

The effect on the model deformation density in the plane of the hydrogen phthalate anion in diaquabis(hydrogen phthalato)copper(II) (17 K data), obtained after refinement with the program *XD* (Koritsanszky *et al.*, 1997), is illustrated in Fig. 4. Remarkably, the double peaks appearing in the cova-

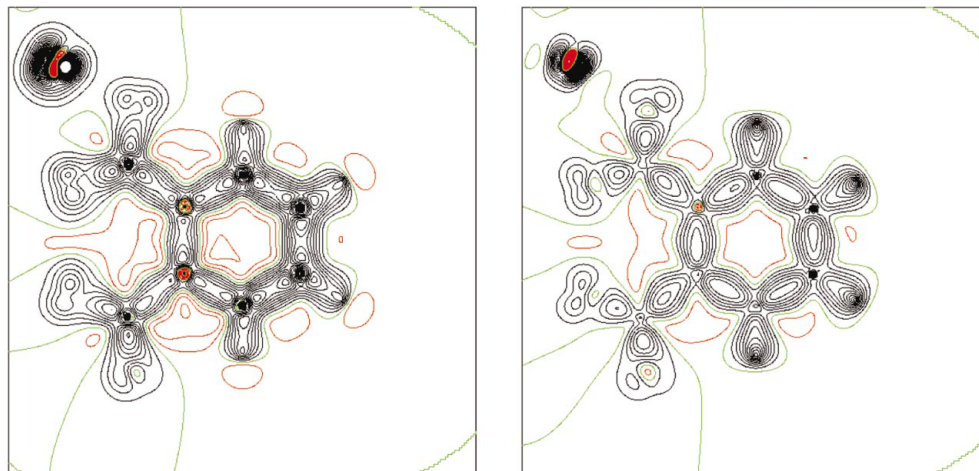


Figure 4

Deformation density maps in the plane of the hydrogen phthalate anion in diaquabis(hydrogen phthalato)copper(II) (17 K data, 0.394 Å). Left: before oblique incident-angle correction. Right: after the correction. Contour interval: 0.10 e Å⁻³; positive contours are black, negative contours red.

lent bonds before correction disappear when the corrected data are used. The 'double bond peak' feature has been observed in other studies, some of which may require re-examination.

5. Concluding remarks

The study reported here concerns one particular type of phosphor. It is clear that a wider examination of the different phosphors being used in crystallographic experiments is needed to ensure the elimination of the systematic errors identified in this study.

We thank Drs Jim Phillips and Roger Durst of Bruker AXS Inc. for supplying the phosphor and for helpful discussions, and Dr Tim Graber of ChemMat CARS at the Advanced Photon Source for assistance with the absorption measurement at the ID-15 beamline at APS. The SUNY X3 beamline at NSLS is supported by the Division of Basic Energy Sciences

of the US Department of Energy (DE-FG02-86ER45231). BLR also thanks CNPq (Brazil) for a post-doctoral fellowship. This research was carried out in part at the National Synchrotron Light Source at Brookhaven National Laboratory, which is supported by the US Department of Energy, Division of Materials Sciences and Division of Chemical Sciences.

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